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                 alerts (SDIs) affected
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NEWS
                 SOLIDSTATE reloaded; updating to resume; current-awareness
                 alerts (SDIs) affected
NEWS
      12 DEC 17
                 CERAB reloaded; updating to resume; current-awareness
                 alerts (SDIs) affected
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     16 JAN 03
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                 February 2005
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     17 JAN 26
                 CA/CAPLUS - Expanded patent coverage to include the Russian
                 Agency for Patents and Trademarks (ROSPATENT)
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=> s (drug delivery) and viscous

2 FILES SEARCHED...

L1 5624 (DRUG DELIVERY) AND VISCOUS

=> s l1 and endocytosis

L2 321 L1 AND ENDOCYTOSIS

=> s 12 and hydrogel

L3 47 L2 AND HYDROGEL

=> s 13 and (apparent viscosity)

L4 7 L3 AND (APPARENT VISCOSITY)

=> d 14 1-7 ibib abs

L4 ANSWER 1 OF 7 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER:

1997:617964 CAPLUS

DOCUMENT NUMBER:

127:268031

TITLE:

Materials and methods for enhancing cellular

internalization

INVENTOR(S):

Edwards, David A.; Deaver, Daniel R.; Langer, Robert

S.

PATENT ASSIGNEE(S):

Penn State Research Foundation, USA; Massachusetts

Institute of Technology

SOURCE:

PCT Int. Appl., 39 pp. CODEN: PIXXD2

Patent

DOCUMENT TYPE:

English

LANGUAGE: En FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE		
						
WO 9732572	A2	19970912	WO 1997-US3276	19970303		
WO 9732572	A3	19971127				

W: AU, CA, JP, KR

RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE

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AU 9720631
                          Α1
                                19970922
                                            AU 1997-20631
                                                                   19970303
                                19981223 EP 1997-908818
     EP 885002
                         A2
                                                                   19970303
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, FI
     US 5985320
                                19991116
                         Α
                                            US 1997-810275
                                                                   19970303
     JP 2000506165
                         T2
                                20000523
                                            JP 1997-531869
                                                                   19970303
PRIORITY APPLN. INFO.:
                                            US 1996-12721P
                                                                P 19960304
                                            WO 1997-US3276
                                                                W
                                                                  19970303
AB
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Compns. and methods for delivering agents across cell membranes are disclosed. The compns. include an agent to be delivered; a viscous material such as a hydrogel, lipogel, or viscous sol; and optionally a carrier that includes a ligand that binds to or interacts with cell surface receptors. The agent to be delivered binds to or otherwise interacts with cell surface receptors; is attached covalently or ionically to a mol. that binds to or interacts with a cell surface receptor; or is associated with the carrier. Agents to be delivered include bioactive compds. and diagnostic agents. The compns. have an apparent viscosity roughly equal to the viscosity of the cytosol in the cell to which the agent is to be delivered. The rate of cellular internalization is higher when the viscosity of the viscous material and that of the cytosol in the cell are approx. the same, relative to when they are not the same. compns. enhance cellular entry of bioactive agents and diagnostic materials when administered vaginally, nasally, rectally, ocularly, orally, or to the respiratory or pulmonary system. Thus, uptake of 125I-labeled transferrin into human K562 erythroleukemia cells by endocytosis from aqueous solns. containing 0.0-1.8% methylcellulose increased slowly with increasing methylcellulose concentration up to 1.25%, then

rapidly up to 1.7%, and decreased rapidly at higher concns.; the apparent viscosity of methylcellulose solns. in the 1.25-1.7% concentration range was similar to that in the K562 cell cytoplasm. Intravaginal administration of 100 μ g leuprolide, a vaginal epithelial LH-RH receptor-binding drug, to sheep in a 1.5% or 1.75% methylcellulose hydrogel resulted in an increase in serum LH concentration

L4 ANSWER 2 OF 7 USPATFULL on STN

ACCESSION NUMBER: 2004:

2004:139366 USPATFULL

TITLE:

Compositions and methods for enhancing receptor-mediated cellular internalization Deaver, Daniel R., Franklin, MA, UNITED STATES

INVENTOR(S):

Deaver, Daniel R., Franklin, MA, UNITED STATES Edwards, David A., Boston, MA, UNITED STATES

PATENT ASSIGNEE(S):

The Penn State Research Foundation (U.S. corporation)

	NUMBER	KIND	DATE	
PATENT INFORMATION: APPLICATION INFO.: RELATED APPLN. INFO.:	US 2004106542 US 2003-717251 Continuation of Apr 2002, GRANTE	A1 A1 Ser. No D, Pat. -412821	20040603 20031119 . US 2002- No. US 66	

NUMBER												D	Α	T	Ε										
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PRIORITY INFORMATION:

US 1998-103117P 19981005 (60)

DOCUMENT TYPE:

Utility

FILE SEGMENT:

APPLICATION

LEGAL REPRESENTATIVE:

PATREA L. PABST, HOLLAND & KNIGHT LLP, SUITE 2000, ONE

ATLANTIC CENTER, 1201 WEST PEACHTREE STREET, N.E.,

ATLANTA, GA, 30309-3400

NUMBER OF CLAIMS:

EXEMPLARY CLAIM:

1

22

NUMBER OF DRAWINGS:

5 Drawing Page(s)

LINE COUNT: 1149

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

Compositions and methods for improving cellular internalization of one or more compounds are disclosed. The compositions include a compound to be delivered and a biocompatible viscous material, such as a hydrogel, lipogel, or highly viscous soluble The composition also include, or are administered in conjunction with, an enhancer in an amount effective to maximize expression of or binding to

receptors and enhance RME of the compound into the cells. This leads to high transport rates of compounds to be delivered across cell membranes, facilitating more efficient delivery of drugs and diagnostic agents. Compositions are applied topically orally, nasally, vaginally, rectally, and ocularly. The enhancer is administered with the composition or separately, either systemically or preferably locally. The compound to be delivered can also be the enhancer.

ANSWER 3 OF 7 USPATFULL on STN L4

ACCESSION NUMBER: 2003:95806 USPATFULL

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TITLE: Process for the preparation of aqueous dispersions of

particles of water-soluble polymers and the particles

obtained

INVENTOR(S): Vanderhoff, John W., Bethlehem, PA, United States

> Lu, Cheng Xun, Somerset, NJ, United States Lee, Clarence C., Lilburn, GA, United States Tsai, Chi-Chun, Lawrenceville, GA, United States

PATENT ASSIGNEE(S): C. R. Bard, Inc., Murray Hill, NJ, United States (U.S.

corporation)

Lehigh University, Bethlehem, PA, United States (U.S.

corporation)

NUMBER KIND DATE -----US 6544503 B1 20030408 US 2000-563037 20000501 (9)

RELATED APPLN. INFO.: Continuation of Ser. No. US 1997-989888, filed on 12

Dec 1997, now patented, Pat. No. US 6214331

Continuation-in-part of Ser. No. US 1996-659770, filed on 6 Jun 1996, now abandoned Continuation-in-part of Ser. No. US 1995-466676, filed on 6 Jun 1995, now

abandoned Utility

DOCUMENT TYPE: FILE SEGMENT: GRANTED

PRIMARY EXAMINER: Page, Thurman K.

ASSISTANT EXAMINER: Nola-Baron, Liliana Di LEGAL REPRESENTATIVE: Kilpatrick Stockton LLP

NUMBER OF CLAIMS: 15 EXEMPLARY CLAIM:

PATENT INFORMATION:

APPLICATION INFO.:

NUMBER OF DRAWINGS: 0 Drawing Figure(s); 0 Drawing Page(s)

LINE COUNT: 3525

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The invention is a process for the preparation of crosslinked water-swellable polymer particles. First, an aqueous polymer solution containing a water-soluble polymer having at least one functional group or charge, is combined with aqueous medium. The aqueous polymer solution is then mixed under moderate agitation with an oil medium and an emulsifier to form an emulsion of droplets of the water-soluble polymer. A crosslinking agent capable of crosslinking the functional groups and/or charges in the water-soluble polymer is then added to the emulsion to form crosslinked water-swellable polymer particles. The invention also includes the particles formed by the process and aqueous dispersions containing the particles which are useful for administering to an individual. The particles of the invention are useful for

implantation, soft tissue augmentation, and scaffolding to promote cell growth.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L4 ANSWER 4 OF 7 USPATFULL on STN

ACCESSION NUMBER: 2002:213430 USPATFULL

TITLE: Compositions and methods for enhancing

receptor-mediated cellular internalization

INVENTOR(S): Deaver, Daniel R., Franklin, MA, UNITED STATES

Edwards, David A., Boston, MA, UNITED STATES

PATENT ASSIGNEE(S): The Penn State Research Foundation (U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 2002114803 A1 20020822 US 6652873 B2 20031125

US 6652873 B2 20031125
APPLICATION INFO.: US 2002-120940 A1 20020410 (10)

RELATED APPLN. INFO.: Continuation of Ser. No. US 1999-412821, filed on 5 Oct

1999, GRANTED, Pat. No. US 6387390

NUMBER DATE

PRIORITY INFORMATION: US 1998-103117P 19981005 (60)

DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: PATREA L. PABST, HOLLAND & KNIGHT LLP, SUITE 2000, ONE

ATLANTIC CENTER, 1201 WEST PEACHTREE STREET, N.E.,

ATLANTA, GA, 30309-3400

NUMBER OF CLAIMS: 22 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 5 Drawing Page(s)

LINE COUNT: 1149

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Compositions and methods for improving cellular internalization of one or more compounds are disclosed. The compositions include a compound to be delivered and a biocompatible **viscous** material, such as a

hydrogel, lipogel, or highly viscous soluble The

composition also include, or are administered in conjunction with, an enhancer in an amount effective to maximize expression of or binding to receptors and enhance RME of the compound into the cells. This leads to high transport rates of compounds to be delivered across cell membranes, facilitating more efficient delivery of drugs and diagnostic agents. Compositions are applied topically orally, nasally, vaginally, rectally, and ocularly. The enhancer is administered with the composition or separately, either systemically or preferably locally. The compound to be delivered can also be the enhancer.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L4 ANSWER 5 OF 7 USPATFULL on STN

INVENTOR (S):

ACCESSION NUMBER: 2002:108620 USPATFULL

TITLE: Compositions and methods for enhancing

receptor-mediated cellular internalization Deaver, Daniel R., Franklin, MA, United States

Edwards, David A., Boston, MA, United States

PATENT ASSIGNEE(S): The Penn State Research Foundation, University Park,

PA, United States (U.S. corporation)

 NUMBER DATE

PRIORITY INFORMATION: US 1998-103117P 19981005 (60)

DOCUMENT TYPE: Utility FILE SEGMENT: GRANTED

PRIMARY EXAMINER: Azpuru, Carlos A.
LEGAL REPRESENTATIVE: Holland & Knight LLP

NUMBER OF CLAIMS: 22 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 10 Drawing Figure(s); 5 Drawing Page(s)

LINE COUNT: 1185

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Compositions and methods for improving cellular internalization of one or more compounds are disclosed. The compositions include a compound to be delivered and a biocompatible viscous material, such as a hydrogel, lipogel, or highly viscous soluble The composition also include, or are administered in conjunction with, an

composition also include, or are administered in conjunction with, an enhancer in an amount effective to maximize expression of or binding to receptors and enhance RME of the compound into the cells. This leads to high transport rates of compounds to be delivered across cell membranes, facilitating more efficient delivery of drugs and diagnostic agents. Compositions are applied topically orally, nasally, vaginally, rectally, and ocularly. The enhancer is administered with the composition or separately, either systemically or preferably locally. The compound to be delivered can also be the enhancer.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L4 ANSWER 6 OF 7 USPATFULL on STN

ACCESSION NUMBER: 2001:51555 USPATFULL

TITLE: Process for the prepa

Process for the preparation of aqueous dispersions of particles of water-soluble polymers and the particles

obtained

INVENTOR(S): Vanderhoff, John W., Bethlehem, PA, United States

Lu, Cheng Xun, Somerset, NJ, United States Lee, Clarence C., Lilburn, GA, United States Tsai, Chi-Chun, Lawrenceville, GA, United States

PATENT ASSIGNEE(S): C. R. Bard, Inc., Murray Hill, NJ, United States (U.S.

corporation)

Lehigh University, Bethlehem, PA, United States (U.S.

Continuation-in-part of Ser. No. US 1996-659770, filed on 6 Jun 1996, now abandoned Continuation-in-part of

corporation)

NUMBER KIND DATE
-----US 6214331 B1 20010410
US 1997-989888 19971212 (8)

Ser. No. US 1995-466676, filed on 6 Jun 1995, now abandoned

PATENT INFORMATION: APPLICATION INFO.:

RELATED APPLN. INFO.:

DOCUMENT TYPE:

FILE SEGMENT: Granted
PRIMARY EXAMINER: Kulkosky, Peter F.

Utility

LEGAL REPRESENTATIVE: Kulkosky, Peter F.

Kulkosky, Peter F.

Kilpatrick Stockton LLP

NUMBER OF CLAIMS: 29
EXEMPLARY CLAIM: 1
LINE COUNT: 3840

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The invention is a process for the preparation of crosslinked water-swellable polymer particles. First, an aqueous polymer solution containing a water-soluble polymer having at least one functional group or charge, is combined with aqueous medium. The aqueous polymer solution is then mixed under moderate agitation with an oil medium and an emulsifier to form an emulsion of droplets of the water-soluble polymer.

A crosslinking agent capable of crosslinking the functional groups and/or charges in the water-soluble polymer is then added to the emulsion to form crosslinked water-swellable polymer particles. The invention also includes the particles formed by the process and aqueous dispersions containing the particles which are useful for administering to an individual. The particles of the invention are useful for implantation, soft tissue augmentation, and scaffolding to promote cell growth.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

ANSWER 7 OF 7 USPATFULL on STN

ACCESSION NUMBER: 1999:146020 USPATFULL

TITLE: Materials and methods for enhancing cellular

internalization

INVENTOR(S): Edwards, David A., State College, PA, United States

Deaver, Daniel R., Port Matilda, PA, United States

Langer, Robert S., Newton, MA, United States

PATENT ASSIGNEE(S): The Penn State Research Foundation, University Park,

PA, United States (U.S. corporation)

NUMBER KIND DATE -----ÚS 5985320 19991116 US 1997-810275 19970303 (8)

NUMBER DATE

-----PRIORITY INFORMATION: US 1996-12721P 19960304 (60)

DOCUMENT TYPE: Utility

FILE SEGMENT: Granted

PRIMARY EXAMINER: Kishore, Gollamudi S. LEGAL REPRESENTATIVE: Monahan, Thomas J.

NUMBER OF CLAIMS: 33 EXEMPLARY CLAIM:

PATENT INFORMATION:

APPLICATION INFO.:

NUMBER OF DRAWINGS: 7 Drawing Figure(s); 7 Drawing Page(s)

LINE COUNT: 991

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

Compositions and methods for delivering agents across cell membranes are disclosed. The compositions include an agent to be delivered, a viscous material, such as a hydrogel, lipogel or viscous sol, and, optionally, a carrier that includes a ligand that binds to or interacts with cell surface receptors. The agent to be delivered binds to or otherwise interacts with cell surface receptors, is attached, either covalently or ionically to a molecule that binds to or interacts with a cell surface receptor, or is associated with the carrier. Agents to be delivered include bioactive compounds and diagnostic agents. The compositions have an apparent viscosity roughly equal to the viscosity of the cytosol in the cell to which the agent is to be delivered. The rate of cellular internalization is higher when the viscosity of the viscous material and that of the cytosol in the cell are approximately the same, relative to when they are not the same. The compositions enhance cellular entry of bioactive agents and diagnostic materials when administered vaginally, nasally, rectally ocularly, orally, or to the respiratory or pulmonary system.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.